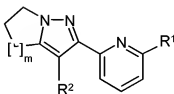


**AMENDMENT TO THE CLAIMS**

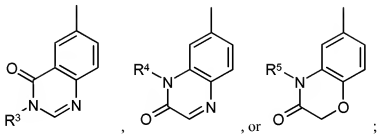
1. (original) A compound of the formula:



Formula I

wherein R<sup>1</sup> is hydrogen or (C<sub>1</sub>-C<sub>6</sub>) alkyl;

R<sup>2</sup> is selected from the group consisting of:



R<sup>3</sup> is (C<sub>1</sub>-C<sub>6</sub>)alkyl, or (CH<sub>2</sub>)<sub>n</sub>X;

R<sup>4</sup> is (C<sub>1</sub>-C<sub>6</sub>)alkyl, or (CH<sub>2</sub>)<sub>n</sub>X;

R<sup>5</sup> is hydrogen, (C<sub>1</sub>-C<sub>6</sub>)alkyl, or (CH<sub>2</sub>)<sub>n</sub>X;

X is selected from the group consisting of a halogen, NR<sup>a</sup>R<sup>b</sup>,

N-morpholino, N-piperidine, N-pyrrolidine, or N-azepane;

n is an integer from 1-4;

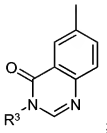
m is an integer from 1-3;

R<sup>a</sup> and R<sup>b</sup> are each independently hydrogen or (C<sub>1</sub>-C<sub>6</sub>)alkyl;  
and the pharmaceutically acceptable salts thereof.

2. (original) A compound according to Claim 1 wherein:

R<sup>1</sup> is hydrogen or (C<sub>1</sub>-C<sub>6</sub>) alkyl;

R<sup>2</sup> is



R<sup>3</sup> is (C<sub>1</sub>-C<sub>6</sub>)alkyl, or (CH<sub>2</sub>)<sub>n</sub>X;

X is selected from the group consisting of a halogen, NR<sup>a</sup>R<sup>b</sup>,

N-morpholino, N-piperidine, N-pyrrolidine, or N-azepane;

n is an integer from 1-4;

m is an integer from 1-3;

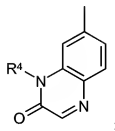
R<sup>a</sup> and R<sup>b</sup> are each independently hydrogen or (C<sub>1</sub>-C<sub>6</sub>)alkyl;

and the pharmaceutically acceptable salts thereof.

3. (original) A compound according to Claim 1 wherein:

R<sup>1</sup> is hydrogen or (C<sub>1</sub>-C<sub>6</sub>) alkyl;

R<sup>2</sup> is



R<sup>4</sup> is (C<sub>1</sub>-C<sub>6</sub>)alkyl, or (CH<sub>2</sub>)<sub>n</sub>X;

X is selected from the group consisting of a halogen, NR<sup>a</sup>R<sup>b</sup>,

N-morpholino, N-piperidine, N-pyrrolidine, or N-azepane;

n is an integer from 1-4;

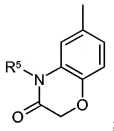
m is an integer from 1-3;

R<sup>a</sup> and R<sup>b</sup> are each independently hydrogen or (C<sub>1</sub>-C<sub>6</sub>)alkyl;  
and the pharmaceutically acceptable salts thereof.

4. (original) A compound according to Claim 1 wherein:

R<sup>1</sup> is hydrogen or (C<sub>1</sub>-C<sub>6</sub>) alkyl;

R<sup>2</sup> is



R<sup>5</sup> is hydrogen, (C<sub>1</sub>-C<sub>6</sub>)alkyl, or (CH<sub>2</sub>)<sub>n</sub>X;

X is selected from the group consisting of a halogen, NR<sup>a</sup>R<sup>b</sup>,

N-morpholino, N-piperidine, N-pyrrolidine, or N-azepane;

n is an integer from 1-4;

m is an integer from 1-3;

R<sup>a</sup> and R<sup>b</sup> are each independently hydrogen or (C<sub>1</sub>-C<sub>6</sub>)alkyl;

and the pharmaceutically acceptable salts thereof.

5. (currently amended) A compound according to ~~Claims 1 to 4~~ Claim 1 wherein m is 1.

6. (currently amended) A compound according to ~~Claims 1 to 5~~ Claim 1 wherein R<sup>3</sup> is methyl.

7. (currently amended) A compound according to ~~Claims 1 to 6~~ Claim 1 wherein R<sup>1</sup> is methyl.

8. (currently amended) A pharmaceutical formulation comprising a compound according to ~~any one of Claims 1 to 7~~ Claim 1 in combination with a pharmaceutically acceptable diluent, excipient or carrier.

9. (currently amended) A method of treating susceptible neoplasms in a patient in need thereof which comprises administering to said patient a therapeutically effective amount of a compound according to ~~any one of Claims 1 to 7~~ Claim 1.

10. (currently amended) A method of treating fibrosis in a patient in need thereof which comprises administering to said patient a therapeutically effective amount of a compound according to ~~any one of Claims 1 to 7~~ Claim 1.